

IN THE CLAIMS:

1-4. (Cancelled)

5. (Currently Amended) A composition comprising:
a cyclodextrin-containing polymer,
a therapeutic agent, and
a complexing agent, comprising:

at least one ~~host/guest~~ moiety at a ~~terminus of the complexing agent~~ that forms an inclusion complex with a ~~host/guest~~ moiety of said cyclodextrin-containing polymer, wherein the guest moiety is selected from adamantyl, naphthyl, cholesterol, and combinations thereof, and

at least one polymer portion that increases solubility and/or imparts stabilization relative to a composition of the cyclodextrin-containing polymer and therapeutic agent alone;

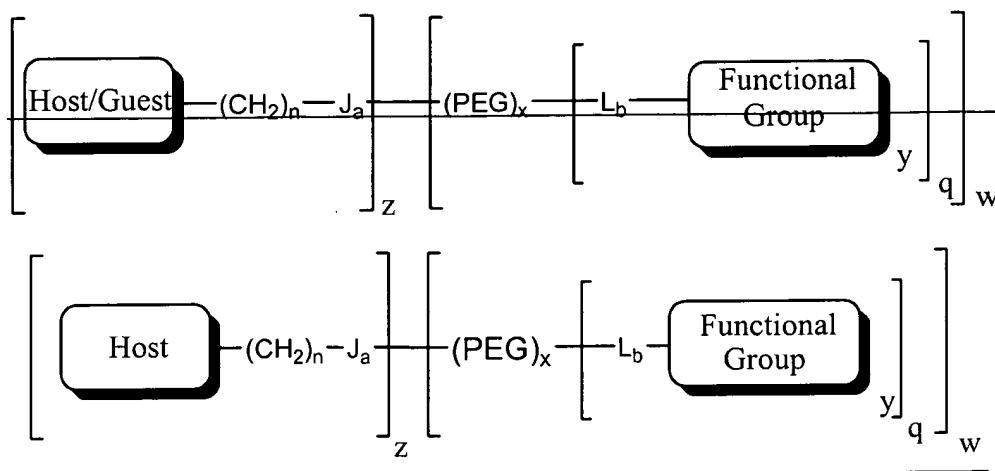
wherein the cyclodextrin-containing polymer, the therapeutic agent, and the complexing agent are separate molecules.

6. (Previously Presented) A composition of claim 5, wherein said therapeutic agent is selected from an antibiotic, a steroid, a polynucleotide, small molecule pharmaceutical, a virus, a plasmid, a peptide, a peptide fragment, a chelating agent, a biologically active macromolecule, and mixtures thereof.

7. (Original) A composition of claim 6, wherein said therapeutic agent is a polynucleotide.

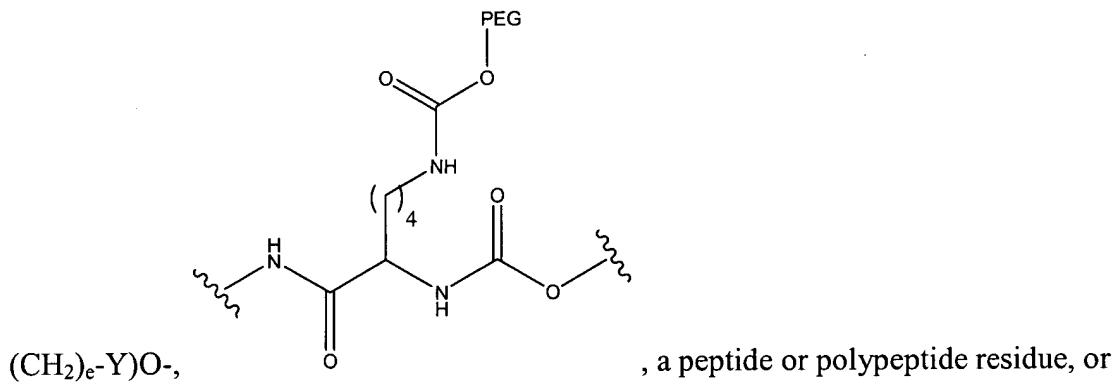
8-11. (Cancelled)

12. (Currently Amended) A composition of claim 5, wherein the complexing agent is a compound of the formula:



wherein

J is $-\text{NH}-$, $-\text{C}(=\text{O})\text{NH}-\text{CH}_2-$, $-\text{NH}-\text{C}(=\text{O})-(\text{CH}_2)_d-$, $-\text{CH}_2\text{SS}-$, $-\text{C}(=\text{O})\text{O}-(\text{CH}_2)_e-\text{O}-\text{P}(=\text{O})(\text{O}-$



Y is an additional host-guest functionality;

R^1 is $-(\text{CH}_2)-\text{CO}_2\text{H}$, an ester or salt thereof; or $-(\text{CH}_2)_a-\text{CONH}_2$;

PEG is $-\text{O}(\text{CH}_2\text{CH}_2\text{O})_z-$, where z varies from 2 to 500;

L is H, -NH, -NH-(C=O)-(CH₂)_e-(C=O)-CH₂-, -S(=O)₂-HC=CH-, -SS-, -C(=O)O-, or a carbohydrate residue;

a is 0 or 1;

b is 0 or 1;

d ranges from 0 to 6;

e ranges from 1 to 6;

n ranges from 0 to 6;

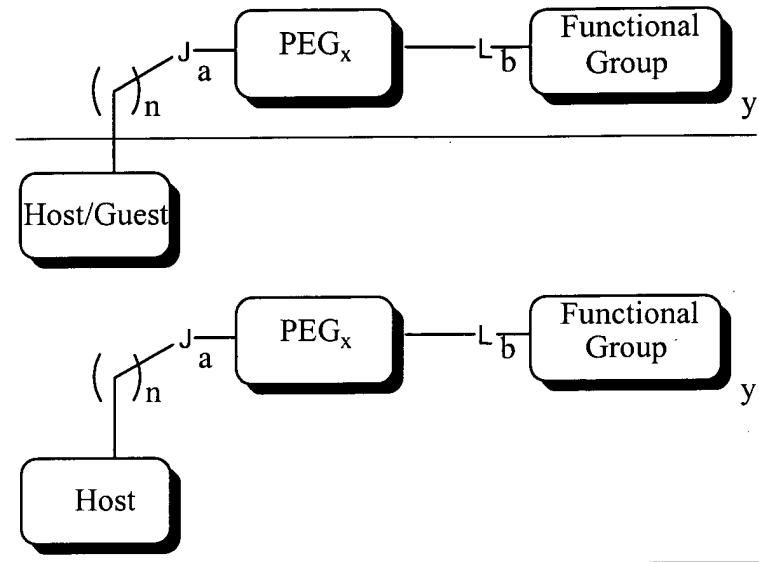
q ranges from 1 to 5;

w ranges from 1 to 5;

y is 1; and

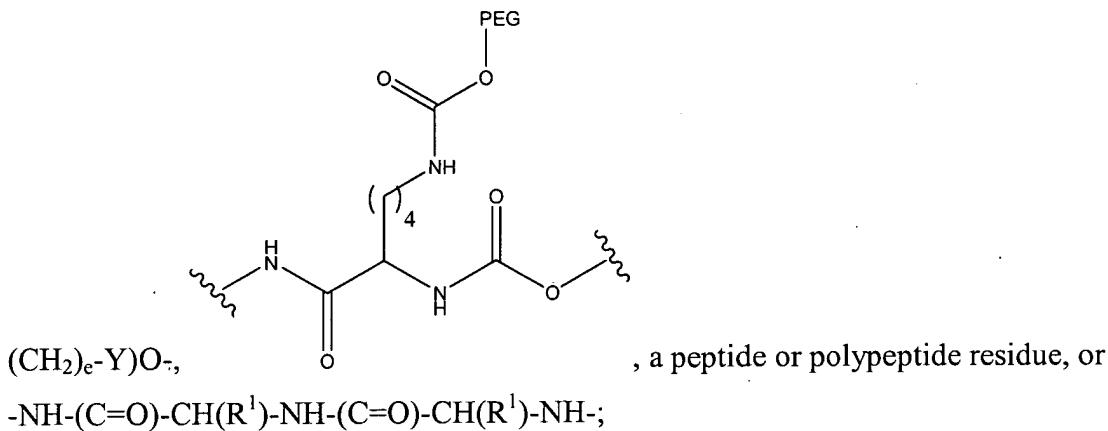
x is 0 or 1.

13. (Currently Amended) A composition of claim 5, wherein the complexing agent is a compound of the formula:



wherein

J is $-\text{NH}-$, $-\text{C}(=\text{O})\text{NH}-\text{CH}_2-$, $-\text{NH}-\text{C}(=\text{O})-(\text{CH}_2)_d-$, $-\text{CH}_2\text{SS}-$, $-\text{C}(=\text{O})\text{O}-(\text{CH}_2)_e-\text{O}-\text{P}(=\text{O})(\text{O}-$



$-\text{NH}-(\text{C}= \text{O})-\text{CH}(\text{R}^1)-\text{NH}-(\text{C}= \text{O})-\text{CH}(\text{R}^1)-\text{NH}-$;

Y is an additional host-guest functionality;

R¹ is $-(\text{CH}_2)-\text{CO}_2\text{H}$, an ester or salt thereof; or $-(\text{CH}_2)_a-\text{CONH}_2$;

PEG is $-\text{O}(\text{CH}_2\text{CH}_2\text{O})_z-$, where z varies from 2 to 500;

L is H, -NH, -NH-(C=O)-(CH₂)_e-(C=O)-CH₂-, -S(=O)₂-HC=CH-, -SS-, -C(=O)O-, or a carbohydrate residue;

a is 0 or 1;

b is 0 or 1;
d ranges from 0 to 6;
e ranges from 1 to 6;
n ranges from 0 to 6;
y is 1; and
x is 0 or 1.

14. (Previously Presented) A composition of claim 5, wherein the complexing agent further comprises a group selected from a ligand, a nuclear localization signal, an endosomal release peptide, an endosomal release polymer, or a membrane permeabilization agent.

15. (Previously Presented) A composition of claim 5, wherein the polymer portion increases the solubility of the composition under biological conditions relative to a composition of the cyclodextrin-containing polymer and therapeutic agent alone.

16. (Previously Presented) A composition of claim 5, wherein the polymer portion stabilizes the composition under biological conditions relative to a composition of the cyclodextrin-containing polymer and therapeutic agent alone.

17. (Previously Presented) A composition of claim 5, wherein the complexing agent further comprises a therapeutic agent reversibly bound to the complexing agent.

18. (Previously Presented) A composition of claim 5, wherein the complexing agent further comprises a spacer group.

19-22. (Cancelled)

23. (Previously Presented) A composition of claim 5, wherein at least one polymer portion of the complexing agent comprises PEG or derivatives thereof.

24-26. (Cancelled)

27. (Previously Presented) A composition of claim 5, wherein the cyclodextrin-containing polymer comprises one or more cyclodextrins in side chains of the cyclodextrin-containing polymer.

28. (Previously Presented) A composition of claim 5, wherein the cyclodextrin-containing polymer comprises a linear cyclodextrin-containing polymer wherein cyclodextrin moieties are present in the backbone of the polymer.

29. (Cancelled)

30. (Currently Amended) A composition comprising:

a cyclodextrin-containing polymer,

a therapeutic agent, and

a complexing agent, comprising:

at least one functional group,

at least one ~~host/guest~~ moiety ~~at a terminus of the complexing agent~~ that forms an inclusion complex with a ~~host/guest~~ moiety of said cyclodextrin-containing polymer, wherein the guest moiety is selected from adamantyl, naphthyl, cholesterol, and combinations thereof, and

at least one polymeric spacer group;

wherein the cyclodextrin-containing polymer, the therapeutic agent, and the complexing agent are separate molecules.

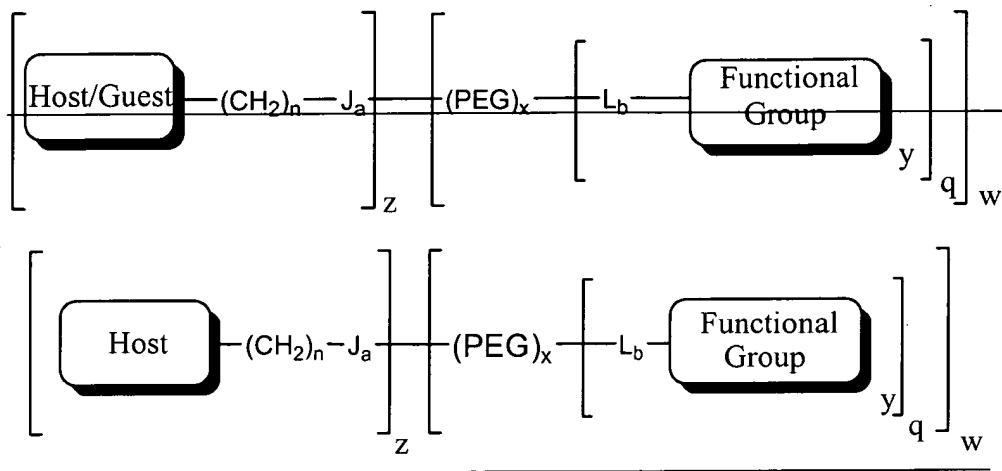
31. (Previously Presented) A composition of claim 30, wherein said therapeutic agent is selected from an antibiotic, a steroid, a polynucleotide, small molecule pharmaceutical, a virus, a plasmid, a peptide, a peptide fragment, a chelating agent, a biologically active macromolecule, and mixtures thereof.

32. (Previously Presented) A composition of claim 31, wherein said therapeutic agent is a polynucleotide.

33. (Cancelled)

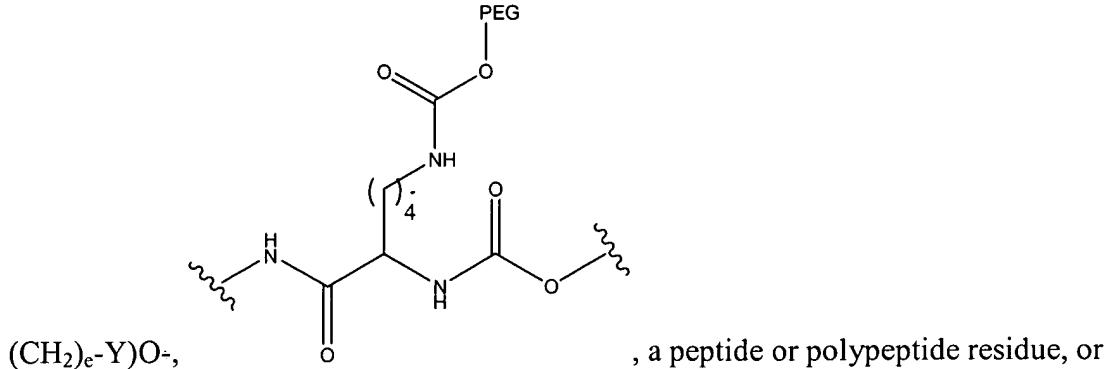
34. (Previously Presented) A composition of claim 30, wherein at least one spacer group of the complexing agent comprises PEG or derivatives thereof.

35. (Currently Amended) A composition of claim 34, wherein the complexing agent is a compound of the formula:



wherein

J is $-\text{NH-}$, $-\text{C(=O)NH-CH}_2\text{d-}$, $-\text{NH-C(=O)-(CH}_2\text{d-}$, $-\text{CH}_2\text{SS-}$, $-\text{C(=O)O-(CH}_2\text{e-O-P(=O)(O-CH}_2\text{e-YO-}$,



Y is an additional host-guest functionality;

R^1 is $-(\text{CH}_2)\text{-CO}_2\text{H}$, an ester or salt thereof; or $-(\text{CH}_2)_a\text{-CONH}_2$;

PEG is $-\text{O}(\text{CH}_2\text{CH}_2\text{O})_z-$, where z varies from 2 to 500;

L is H, $-\text{NH-}$, $-\text{NH-C(=O)-(CH}_2\text{e-C(=O)-CH}_2\text{-}$, $-\text{S(=O)}_2\text{-HC=CH-}$, $-\text{SS-}$, $-\text{C(=O)O-}$, or a carbohydrate residue;

a is 0 or 1;

b is 0 or 1;

d ranges from 0 to 6;

e ranges from 1 to 6;

n ranges from 0 to 6;

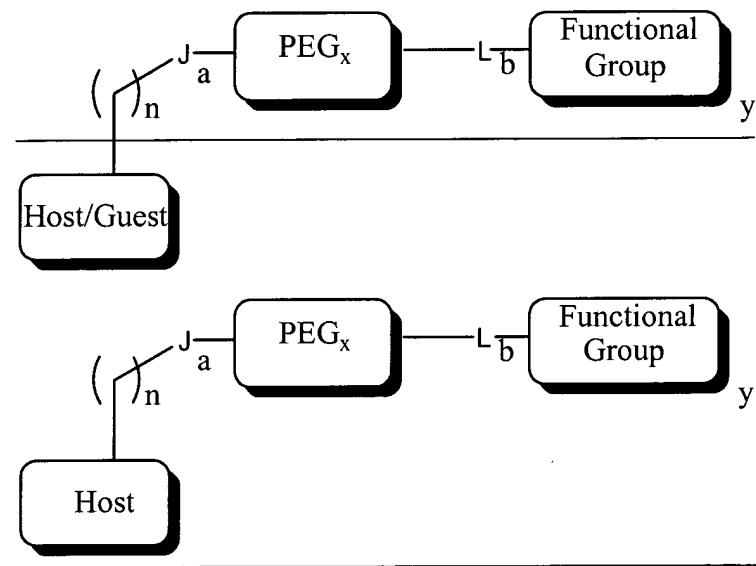
q ranges from 1 to 5;

w ranges from 1 to 5;

y is 1; and

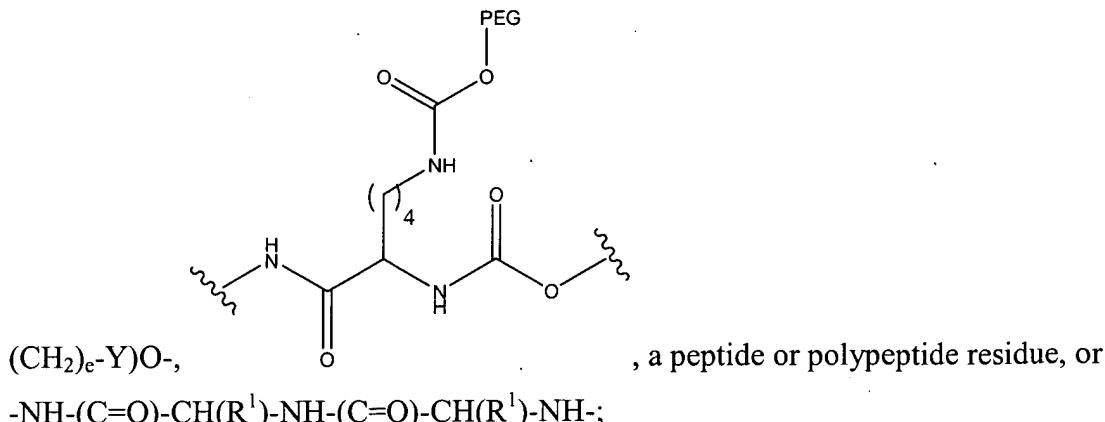
x is 1.

36. (Currently Amended) A composition of claim 34, wherein the complexing agent is a compound of the formula:



wherein

J is $-\text{NH}-$, $-\text{C}(\text{=O})\text{NH}-\text{CH}_2-$, $-\text{NH}-\text{C}(\text{=O})-(\text{CH}_2)_d-$, $-\text{CH}_2\text{SS}-$, $-\text{C}(\text{=O})\text{O}-(\text{CH}_2)_e-\text{O}-\text{P}(\text{=O})(\text{O}-$



Y is an additional host-guest functionality;

R¹ is $-(\text{CH}_2)-\text{CO}_2\text{H}$, an ester or salt thereof; or $-(\text{CH}_2)_a-\text{CONH}_2$;

PEG is $-\text{O}(\text{CH}_2\text{CH}_2\text{O})_z-$, where z varies from 2 to 500;
L is H, $-\text{NH}-$, $-\text{NH}-(\text{C}=\text{O})-(\text{CH}_2)_e-(\text{C}=\text{O})-\text{CH}_2-$, $-\text{S}(=\text{O})_2-\text{HC}=\text{CH}-$, $-\text{SS}-$, $-\text{C}(=\text{O})\text{O}-$, or a carbohydrate residue;

a is 0 or 1;

b is 0 or 1;

d ranges from 0 to 6;

e ranges from 1 to 6;

n ranges from 0 to 6;

y is 1; and

x is 1.

37. (Previously Presented) A composition of claim 30, wherein at least one functional group includes a group selected from a ligand, a nuclear localization signal, an endosomal release peptide, an endosomal release polymer, or a membrane permeabilization agent.

38. (Previously Presented) A composition of claim 30, wherein at least one functional group includes a moiety that increases the solubility of the composition under biological conditions relative to a composition of the cyclodextrin-containing polymer and therapeutic agent alone.

39. (Previously Presented) A composition of claim 30, wherein at least one functional group includes a moiety that stabilizes the composition under biological conditions relative to a composition of the cyclodextrin-containing polymer and therapeutic agent alone.

40. (Previously Presented) A composition of claim 30, wherein at least one functional group includes a therapeutic agent reversibly bound to the complexing agent.

41. (Previously Presented) A composition of claim 30, wherein the cyclodextrin-containing polymer comprises one or more cyclodextrins in side chains of the cyclodextrin-containing polymer.

42. (Previously Presented) A composition of claim 30, wherein the cyclodextrin-containing polymer comprises a linear cyclodextrin-containing polymer wherein cyclodextrin moieties are present in the backbone of the polymer.